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22850	7590 04/20/2004		EXAMINER	
OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT, P.C. 1940 DUKE STREET			MORAMED	, ABDEL A
ALEXANDRIA, VA 22314			ART UNIT	PAPER NUMBER
			1653	
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Please find below and/or attached an Office communication concerning this application or proceeding.

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OBLON, SPIVAK, N	ACCLELLAND
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DETAILED ACTION

ACKNOWLEDGMENT OF THE SUPPLEMENTAL AND APPEAL BRIEFS, AMENDMENT, STATUS OF THE APPLICATION AND CLAIMS

1. The request for reconsideration of Supplemental Appeal Brief filed 1/30/04, Supplemental Appeal Brief and Supplemental amendment filed 5/12/03 and Appeal Brief filed 3/31/03 are acknowledged, entered and considered. In view of Applicant's request (Supplemental Amendment filed 5/12/03) claims 36 and 47 have been canceled and claims 53 and 54 have been added. Thus, claims 31-35, 37-46 and 48-54 are now pending in the application. The rejections under 35 U.S.C. 103(a) over the prior art of record and 35 U.S.C. 112, second paragraph are withdrawn in view of Applicant's amendment and arguments filed 5/12/03. The Finality of the previous Office action is withdrawn in view of the following new ground of rejection:

CLAIMS REJECTION-35 U.S.C. § 103(a)

- 2. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:
- (a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein

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were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 GFR 0.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(f) or (g) prior art under 35 U.S.C. 103(a).

Claims 31-35, 37-46 and 48-54 are rejected under 35 U.S.C. 103(a) as being unpatentable over Daugherty et al. (The Journal of Biological Chemistry, Vol. 273, No. 51, pp. 33961-33971, December 18, 1998) taken with Kitamura et al. (Macromol. Rapid Commun., Vol. 20, pp. 612-615, 1999) and Larsen et al. (Carbohydrate Research, Vol. 309, pp. 153-159, 1998).

The reference of Daugherty et al. discloses the application of an artificial chaperone refolding methods to porcine heart citrate synthase (CS), carbonic anhydrase B (CAB) and lysozyme by combining cyclic saccharide such as cyclodextrin and detergents such as POE(10)L, Triton X-100, SDS, etc. i.e., nonionic and ionic detergents (See e.g. pages 33961 and 33963-33964) as directed to claims 31-35, 37, 38 and 53. It is noted that the reference does not recite cyclic saccharide cycloamylose as currently claimed in claims 31 and 35; however, the reference recites cyclodextrin which is defined as a high molecular weight cyclic σ -1,4-glucan which is referred as cycloamylose and generally called σ , ρ and ρ cyclodextrin, respectively. Thus, it is known in the art that cyclodextrin is the same as cycloamylose. For support, See e.g., particularly the reference of Machida et al. at page 131 (Machida et al. FEBS Letters, 486 (2000) 131-135) which was attached as pertinent art in the previous Office action

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mailed on 8/13/02 as Paper No.6. Thus, the reference of Daugherty et al. discloses the combination of cyclic saccharide cycloamylose and detergents such as polyoxyethylenic detergent or ionic detergent such as SDS are useful as artificial chaperone.

The reference also teaches the use of the artificial chaperone refolding method to porcine heart citrate synthase wherein the folded protein has an α-helical structure or β-sheet structure or has an intramolecular S-S bond comprising of two steps, wherein the first step is to prevent aggregation of protein molecules by the formulation of protein-detergent complexes, in which hydrophobic regions of non-native protein molecules are shielded by detergent. The second step is the addition of cyclodextrin (cycloamylose) which initiates folding by stripping detergent from the protein-detergent complexes to facilitate the proper folding-of protein into a correct higher-order structure with activity (See e.g. pages 33961, 33963-33964, 33966-33971 and Scheme I) as directed to claims 39-46, 48-52 and 54.

Although, the primary reference of Daugherty et al. clearly teaches the combination of cyclic saccharide cycloamylose and detergents such as polyoxyethylenic or ionic detergents is useful as an artificial chaperone and method for diluting the denaturant thereof and method of refolding denatured protein into native state having an activity by using the artificial chaperone as claimed in claims 31-35, 37-46 and 48-54. However, the reference differs from claims 31-35, 37-46 and 48-54 in not teaching the use of an artificial chaperone kit and the recitation of specific polymerization degree of cyclic saccharide cycloamylose. Nevertheless, the secondary reference of Kitamura et al. states that one of the remarkable properties of common cyclodextrins (α-, β-, and γ-

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CD) is their ability to include small molecules (artificial chaperones) within their cyclic structure to form inclusion compounds. Most recently, large cyclodextrins with degrees of polymerization (DP) from 17 to several hundreds have been found as the result of the action of recombinant potato D-enzyme. These large CDs may have the potential to function as host molecules for a variety of organic reagents and iodine in a manner, which is different from the common cyclodextrins (See e.g., page 612, first paragraph, left column). Hence, describing cyclic saccharide cycloamylose (i.e., cyclodextrin) having a polymerization degree from 17 to several hundreds, and as such overlaps with the claimed ranges of polymerization degrees of claims 31, 33-35, 37-39, 41, 42, 46, 48. and 49. Further, the secondary reference of Larsen et al. on page 154, second paragraph, left column states that the large interest in cyclodextrins is based on their ability to form inclusion complexes with wide range of molecules. α -, β -, and γ -CD are capable to complex molecules ranging from gases to proteins and other biopolymers. CD is able to modify the physicochemical properties of the guest molecules by increasing their solubility and stability, as well as by modifying their reactivity. They have found numerous applications in the agricultural, food, chemical, and pharmaceutical industries. Furthermore, CD has been shown to be valuable as selectivity reagents for the resolution of structural, positional and stereoisomers in analytical chemistry. On page 156, the reference concludes by stating that using this method we have found that large CD have the ability to form complexes similar to the small CD (α -, β , and γ -CD). θ -CD gave the highest inclusion complex formation

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constant among the larger CD. Thus, the secondary references discuss information on larger CD and their inclusion complex forming properties.

Therefore, it would have been obvious to one of ordinary skill in the art to apply the teachings of the secondary references of using different cyclodextrins (i.e., α -, β , and γ -CD) and their inclusion complex properties to the primary references teachings of protein refolding with cycloamylose because such features are known or suggested in the art, as seen in the secondary references, and including such features into the methods of the primary reference would have been obvious to one of ordinary skill in the art to obtain the known and recognized functions and advantages of increasing the solubility, stability and modifying the reactivity of the molecules with which they can from inclusion complexes thereof.

In regard to the kit, the primary reference discloses an artificial chaperon formulation, however, from the cited references, it is conventional and within the ordinary skill in the art based upon the teachings of the combined references to have such kits/compositions as set forth in claims 31-35, 37, 38 and 53 since the combined references teach using these compositions together in the same formulations that would have been found in the claimed compositions and/or kits to formulate compositions into a kit format because the claimed kit is tailored for use in the claimed artificial chaperone kit formulation comprising the composition claimed. Hence, it would have been obvious to package the composition required for the method into kit format of the well-known commercial expediency of doing so.

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Therefore, in view of the above and in view of the combined teachings of the prior art, one of ordinary skill in the art would have been motivated at the time the invention was made to employ an artificial chaperone kit comprising cyclic saccharide cycloamylose and polyoxyethylenic detergent or cyclic saccharide cycloamylose and ionic detergent and to a method for diluting the denaturant making the protein a denatured state by adding a specific detergent to a denatured protein, and preventing protein molecules from aggregation, thereafter adding cyclic saccharide cycloamylose, utilizing the inclusion ability thereof to strip detergent, accelerating the proper folding of protein into a correct higher-order structure activity, absence of sufficient objective factual evidence or unexpected results to the contrary.

CONCLUSION AND FUTURE CORRESPONDENCE

6. No claim is allowed.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Abdel A. Mohamed whose telephone number is (571) 272-0955. The examiner can normally be reached on Monday through Friday from 7:30 a.m. to 5:00 p.m. The examiner can also be reached on alternate Fridays.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Christopher Low, can be reached on (571) 272-0951. The appropriate fax phone number for the organization where this application or proceeding is assigned arte (703) 3872-9306 for regular communications and (703) 305-7401 for After Final communications.

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Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (571) 272-1600.

AM Mohamed/AAM

April 16, 2004

Chmis lop fur S. J. Low CHRISTOPHER S. F. LOW SUPERVISORY PATENT EXAMINER TECHNOLOGY CENTER 1800